

Appl. No 10/561,162
Reply to Office Action of April 16, 2008

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AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

1. (Withdrawn) A method for preventing or attenuating atrial fibrillation (AF) promotion by atrial tachycardia in a subject comprising the administration of a therapeutically effective amount of a HMG-CoA reductase inhibitor.
2. (Withdrawn) A method as defined in claim 1, wherein said HMG-CoA reductase inhibitor is effective against longer-term atrial tachycardia remodeling.
3. (Original) A method as defined in claim 2, wherein said longer-term is greater than 24 hours.
4. (Withdrawn) A method as defined in claim 2, wherein said HMG-CoA reductase inhibitor is selected from the group consisting of: atorvastatin (Lipitor®), cerivastatin (Baycol®), fluvastatin (Lescol®), lovastatin (Mevacor®, Altocor®), pravastatin (Pravachol®), simvastatin (Zocor®), epistatin, eptastatin, mevinolin, and synvinolin.
5. (Withdrawn) A method as defined in claim 4, wherein said HMG-CoA

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reductase inhibitor is simvastatin (Zocor®).

6. (Withdrawn) A method as defined in claim 4, wherein said HMG-CoA reductase inhibitor is administered in an amount of about 0.1-2 mg/day.

7. (Withdrawn) A method as defined in claim 6, wherein said subject is a mammal.

8. (Withdrawn) A method as defined in claim 7, wherein said mammal is human.

9. (Currently amended) A method of reducing the incidence of preventing atrial fibrillation (AF) by substrate modification comprising the step of administering to a mammal subject in need thereof an amount of a statin drug therapeutically effective for reducing the incidence of atrial fibrillation in the mammal therapeutically effective amount of a statin drug.

10. (Cancelled)

11. (Original) A method as defined in claim 10, wherein said statin drug is simvastatin (Zocor®).

12. (Currently amended) A method as defined in claim 11, wherein said statin drug is administered in an amount of about 0.1-2 mg/day.

13. (Cancelled)

14. (Currently amended) A method as defined in claim 9~~claim 13~~, wherein said mammal is human.

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15. (Withdrawn) A method of attenuating atrial tachypacing (ATP) effects on effective refractory period (ERP) in right atrium (RA) appendage, posterior wall and inferior wall comprising the step of administering to a subject in need thereof a therapeutically effective amount of a statin drug.
16. (Withdrawn) A method as defined in claim 15, wherein said statin drug is chosen from the group consisting of: atorvastatin (Lipitor®), cerivastatin (Baycol®), fluvastatin (Lescol®), lovastatin (Mevacor®, Altocor®), pravastatin (Pravachol®), simvastatin (Zocor®), epistatin, eptastatin, mevinolin, and synvinolin.
17. (Withdrawn) A method as defined in claim 16, wherein said statin drug is simvastatin (Zocor®).
18. (Withdrawn) A method as defined in claim 16, wherein said statin drug is administered in an amount of about 0.1-2 mg/day.
19. (Withdrawn) A method as defined in claim 18, wherein said subject is a mammal.
20. (Withdrawn) A method as defined in claim 19, wherein said mammal is human.
21. (Withdrawn) A method for modulating atrial tachycardia-induced effects on CaV1.2 protein expression, said method comprising the step of administering to a subject an effective amount of a statin drug.
22. (Withdrawn) A method as defined in claim 21, wherein said statin drug

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is chosen from the group consisting of: atorvastatin (Lipitor®), cerivastatin (Baycol®), fluvastatin (Lescol®), lovastatin (Mevacor®, Altocor®), pravastatin (Pravachol®), simvastatin (Zocor®), epistatin, eptastatin, mevinolin, and synvinolin.

23. (Withdrawn) A method as defined in claim 22, wherein said statin drug is simvastatin (Zocor®).

24. (Withdrawn) A method as defined in claim 22, wherein said statin drug is administered in an amount of about 0.1-2 mg/day.

25. (Withdrawn) A method as defined in claim 24, wherein said subject is a mammal.

26. (Withdrawn) A method as defined in claim 25, wherein said mammal is human.